Cleft lip/palate-intestinal malrotation-cardiopathy syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Cleft lip/palate-intestinal malrotation-cardiopathy syndrome. ORPHA:2001

Cleft lip/palate-intestinal malrotation-cardiopathy is a multiple congenital anomaly syndrome characterized by flat face, hypertelorism, flat occiput, upward slanting palpebral fissures, cleft palate, micrognathia, short neck, and severe congenital heart defects. Malrotation of the intestine, bilateral clinodactyly, bilobed tongue, short fourth metatarsals and bifid thumbs may be additionally observed.